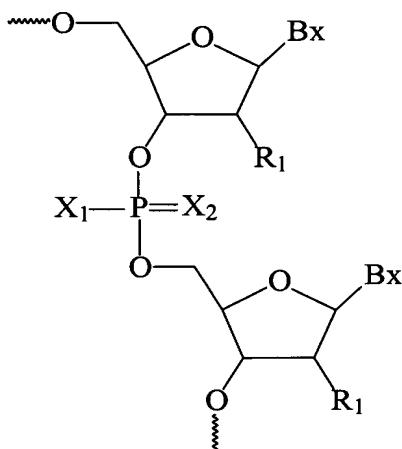


This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Original) A method of preparing an oligomeric compound having at least one moiety of formula:



wherein:

$X_2$  is O or S;

$X_1$  is Pg-O-, Pg-S-,  $C_1$ - $C_{10}$  straight or branched chain alkyl,  $CH_3(CH_2)_{nn}$ -O-,

$R_2R_3N$ - or a group remaining from coupling a chiral auxiliary;

$nn$  is from 0 to 10;

Pg is  $CH_3$ ,  $-CH_2CH_2CN$ ,  $-C(CH_3)(CH_3)-CCl_3$ ,  $-CH_2-CCl_3$ ,  $-CH_2CH=CH_2$ ,

$CH_2CH_2SiCH_3$ , 2-yl-ethyl phenylsulfonate,  $\delta$ -cyanobutenyl, cyano *p*-xylyl, diphenylsilylethyl,

4-nitro-2-yl-ethylbenzene, 2-yl-ethyl-methyl sulfonate, methyl-N-trifluoroacetyl ethyl, acetoxy phenoxy ethyl, or a blocking group;

each  $R_2$  and  $R_3$  is, independently, hydrogen,  $C_1$ - $C_{10}$  alkyl, cycloalkyl or aryl;

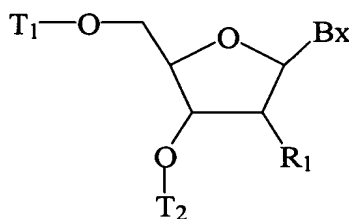
or optionally,  $R_2$  and  $R_3$ , together with the nitrogen atom to which they are attached form a cyclic moiety;

each Bx is, independently, a heterocyclic base moiety; and

each  $R_1$  is, independently, H, a blocked hydroxyl group, or a sugar substituent group;

comprising the steps of:

(a) providing a 5'-O-protected compound of the formula:



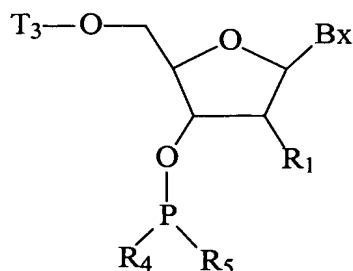
wherein:

$T_1$  is a hydroxyl protecting group; and

$T_2$  is a covalent attachment to a support media, a nucleoside bound to a support media, a nucleotide, an oligonucleoside or an oligonucleotide;

(b) treating said 5'-O-protected compound with a deprotecting reagent for a time and under conditions effective to form a 5'-O-deprotected compound;

(c) coupling said 5'-O-deprotected compound with an activated phosphorus composition of the formula:



wherein:

$T_3$  is a hydroxyl protecting group, a nucleoside, a nucleotide, an oligonucleoside or an oligonucleotide;

$R_4$  is  $N(L_1)L_2$ ;

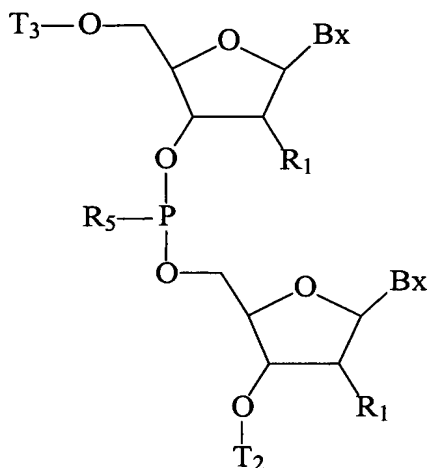
each  $L_1$  and  $L_2$  is, independently,  $C_{1-6}$  straight or branched alkyl, or a  $C_{5-7}$  cyclic aliphatic ring system;

or  $L_1$  and  $L_2$  are joined together to form a 4- to 13-membered heterocyclic ring system including the nitrogen atom to which  $L_1$  and  $L_2$  are attached; and

$R_5$  is  $X_1$ ;

or  $R_4$  and  $R_5$  together with the phosphorus atom to which  $R_4$  and  $R_5$  are attached form a chiral auxiliary;

for a time and under conditions effective to form an extended compound having the formula:



(d) treating said extended compound with a mixture comprising an oxidizing reagent and a capping reagent for a time and under conditions effective to form said oligomeric compound.

2. (Original) The method of claim 1 further comprising treating said oligomeric compound with a reagent for a time and under conditions effective to remove said blocking groups thereby forming a deblocked oligomeric compound.

3. (Original) The method of claim 2 wherein said reagent is effective to cleave the oligomeric compound from the support media.

4. (Original) The method of claim 3 wherein said reagent is aqueous ammonium hydroxide.

5. (Original) The method of claim 2 further comprising treating said oligomeric compound with a further reagent for a time and under conditions effective to cleave the oligomeric compound from the support media.
6. (Original) The method of claim 1 further comprising treating said oligomeric compound with a deprotecting reagent for a time and under conditions effective to deprotect the T<sub>3</sub> hydroxyl protecting group.
7. (Original) The method of claim 1 wherein said mixture comprises from 0.02M to 0.2M oxidizing reagent.
8. (Original) The method of claim 7 wherein said mixture comprises from 0.1M to 0.2M oxidizing reagent.
9. (Original) The method of claim 1 wherein said oxidizing reagent transfers an oxygen atom.
10. (Original) The method of claim 9 wherein said oxidizing reagent is iodine, *m*-chloroperbenzoic acid, iodobenzene diacetate, tetra-*n*-butylammonium periodate, *tert*-butyl hydroperoxide, di-*tert*-butyl hydroperoxide, cumene hydroperoxide, hydrogen peroxide; bis-

trimethylsilyl peroxide; dinitrogen tetroxide, oxone, molecular oxygen, (1*S*)-(+)-(10-camphorsulfonyl)oxaziridine or a peracid.

11. (Original) The method of claim 10 wherein said oxidizing reagent is iodine, *m*-chloroperbenzoic acid, iodobenzene diacetate, *tert*-butyl hydroperoxide, di-*tert*-butyl hydroperoxide, hydrogen peroxide, oxone, molecular oxygen or a peracid.

12. (Original) The method of claim 1 wherein said oxidizing reagent transfers a sulfur atom.

13. (Original) The method of claim 12 wherein said oxidizing reagent is 3-amino-1,2,4-dithiazole-5-thione; 3-ethoxy-1,2,4-dithiazoline-5-one; 1,2,4-dithiazolidine-3,5-dione; 3-methyl-1,2,4-dithiazolin-5-one; or dimethylthiuram disulfide.

14. (Original) The method of claim 13 wherein said oxidizing reagent is dimethylthiuram disulfide.

15. (Original) The method of claim 1 wherein said capping reagent comprises about one part by volume of either acetic anhydride in acetonitrile or tetrahydrofuran; or chloroacetic anhydride in acetonitrile or tetrahydrofuran; added to about one part by volume of

either N-methylimidazole and pyridine in acetonitrile or tetrahydrofuran; or *t*-butylphenoxyacetic anhydride in acetonitrile or tetrahydrofuran.

16. (Original) The method of claim 15 wherein said capping reagent comprises about one part by volume of 20% acetic anhydride in acetonitrile mixed with about one part by volume of a solution having 20% N-methylimidazole, 30% pyridine and 50% acetonitrile.

17. (Original) The method of claim 1 wherein said mixture comprises dimethylthiuram disulfide, acetic anhydride, acetonitrile, N-methyl imidazole and pyridine.

18. (Original) The method of claim 1 wherein said mixture comprises from about 0.05M to 0.2M dimethylthiuram disulfide, about 10% acetic anhydride, about 10% N-methyl imidazole and about 15% pyridine in a suitable solvent.

19. (Original) The method of claim 18 wherein said solvent is acetonitrile, toluene, ethyl acetate, tetrahydrofuran, dichloromethane, dichloroethane, dioxane, dimethylacetamide and dimethylformamide.

20. (Original) The method of claim 1 wherein said coupling of the 5'-O-deprotected compound with the activated phosphorus composition is performed in the presence of an activating agent.

21-78. (Cancelled)